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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.
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08/480, 494

06/07/95

ROESKE

R PPI-007

LAHIVE AND COCKFIELD
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BOSTON MA 02109-1875

18N2/0602

EXAMINER

BORIN, M

ART UNIT

PAPER NUMBER

18

1811

DATE MAILED:

06/02/97

Please find below and/or attached an Office communication concerning this application or proceeding.

Commissioner of Patents and Trademarks

See the attached.



UNITED STATES DEPARTMENT OF COMMERCE
Patent and Trademark Office
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SERIAL NUMBER	FILING DATE	FIRST NAMED APPLICANT	ATTORNEY DOCKET NO.
08/480494	06/07/95	Roeske R.W.	PPI-007

EXAMINER	
Michael L. Borin	
ART UNIT	PAPER NUMBER
1811	18
DATE MAILED:	

Please find below a communication from the EXAMINER in charge of this application

Commissioner of Patents

The communication filed on 5/16/97 is not fully responsive to the communication mailed 3/26/97 for the reason(s) set forth on the attached Notice to Comply With the Sequence Rules or CRF Diskette Problem Report.

Since the response appears to be bona fide, but through an apparent oversight or inadvertence failed to provide a complete response, applicant is required to complete the response within a time limit of one (1) month from the date of this letter or within the time remaining in the response period of the communication mailed [mail date], whichever is the longer. 37 CFR 1.135(c).

NO EXTENSION OF THIS TIME LIMIT MAY BE GRANTED UNDER EITHER 37 C.F.R. 1.136(a) OR (b), BUT THE STATUTORY PERIOD FOR RESPONSE SET IN THE COMMUNICATION MAILED [mail date] MAY BE EXTENDED UP TO A MAXIMUM OF SIX (6) MONTHS UNDER 37 CFR 1.136.

Any inquiry concerning this communication should be directed to Examiner H.A. Gap, Art Unit 1894, whose telephone number is (703) 123-4567.

Any inquiry of a general nature or relating to the status of this application should be directed to the Group receptionist whose telephone number is (703) 308-0196.

CT
CECILIA J. TSANG
SUPERVISORY PATENT EXAMINER
GROUP 1800

**NOTICE TO COMPLY WITH REQUIREMENTS FOR PATENT APPLICATIONS CONTAINING
NUCLEOTIDE SEQUENCE AND/OR AMINO ACID SEQUENCE DISCLOSURES**

The nucleotide and/or amino acid sequence disclosure contained in this application does not comply with the requirements for such a disclosure as set forth in 37 C.F.R. 1.821 - 1.825 for the following reason(s):

- 1. This application clearly fails to comply with the requirements of 37 C.F.R. 1.821-1.825. Applicant's attention is directed to these regulations, published at 1114 OG 29, May 15, 1990 and at 55 FR 18230, May 1, 1990.
- 2. This application does not contain, as a separate part of the disclosure on paper copy, a "Sequence Listing" as required by 37 C.F.R. 1.821(c).
- 3. A copy of the "Sequence Listing" in computer readable form has not been submitted as required by 37 C.F.R. 1.821(e).
- 4. A copy of the "Sequence Listing" in computer readable form has been submitted. However, the content of the computer readable form does not comply with the requirements of 37 C.F.R. 1.822 and/or 1.823, as indicated on the attached copy of the marked -up "Raw Sequence Listing."
- 5. The computer readable form that has been filed with this application has been found to be damaged and/or unreadable as indicated on the attached CRF Diskette Problem Report. A Substitute computer readable form must be submitted as required by 37 C.F.R. 1.825(d).
- 6. The paper copy of the "Sequence Listing" is not the same as the computer readable from of the "Sequence Listing" as required by 37 C.F.R. 1.821(e).
- 7. Other: The sequence listing presented provides the sequence (SEQ ID #1) for the natural LHRH only. The sequence listings for the majority of claimed compounds, subject to Sequence Rules, have not been submitted. Peptides consisting entirely of four or more L-amino acids, whether naturally occurring or not, are subject to the rules. The non-naturally occurring amino acids are represented as Xaa in the Sequence Listing. All amino acid sequences recited in either the specification or the drawings must be followed by a SEQ ID NO. See 37 CFR 1.821(d). With respect to the use of the SEQ ID NOs in the claims, the examiner recommends that the SEQ ID NOs be placed inside of the semicolons immediately following the amino acid sequences. Should this application issue as a patent, there is no guarantee that the printer will print the claims in a format having only one sequence per line, and if the sequences are not so printed it will appear as though the SEQ ID NOs correspond to the following sequences.

Applicant must provide a substitute computer readable form (CRF) copy of the Sequence Listing, a substitute paper copy of the Sequence Listing as well as an amendment directing its entry into the specification, and a statement that the content of the paper and computer readable copies are the same and include no new matter as required by 37 CFR 1.825(a) and (b).

Applicant Must Provide:

- An initial or substitute computer readable form (CRF) copy of the "Sequence Listing".

- An initial or substitute paper copy of the "Sequence Listing", as well as an amendment directing its entry into the specification.
- A statement that the content of the paper and computer readable copies are the same and, where applicable, include no new matter, as required by 37 C.F.R. 1.821(e) or 1.821(f) or 1.821(g) or 1.825(b) or 1.825(d).

For questions regarding compliance to these requirements, please contact:

For Rules Interpretation, call (703) 308-4216
For CRF Submission Help, call (703) 308-4212
For PatentIn software help, call (703) 308-6856

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